

# K210680 AcQMap High Resolution Imaging and Mapping System

Aug 5, 2021  
150 days to decisionK210680 · Product code: **DQK** · Cardiovascular  
Source: <https://www.510kdatabase.net/k210680/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)       |
| Submission type       | Traditional                              |
| Device classification | Computer, Diagnostic, Programmable (DQK) |
| Date received         | Mar 8, 2021                              |
| Decision date         | Aug 5, 2021                              |
| Days to decision      | 150 days                                 |
| Third-party review    | No                                       |
| Combination product   | No                                       |
| PCCP authorized       | No                                       |
| Summary / Statement   | Summary                                  |

**APPLICANT**

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|----------------|---|
| Company        | <b>Acutus Medical, Inc.</b>             |
| Location       | Carlsbad, CA, US                        |
| Contact        | Sindhu Sridhar                          |
| 510(k) history | 24 submissions · 24 cleared · 2017-2023 |

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k210680/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 13, 2026